IN THE UNITED STATES DISTRICT COURT FOR THE DISTRICT OF DELAWARE

AVENTIS PHARMACEUTICALS INC. and SANOFI-AVENTIS US LLC,))
Plaintiffs,)))
v.)
BARR LABORATORIES, INC.)
Defendant.)
)

NOTICE OF DEPOSITION PURSUANT TO FED. R. CIV. P. 30(b)(6)

PLEASE TAKE NOTICE that, pursuant to Rule 30(b)(6) of the Federal Rules of Civil Procedure, Defendant Barr Laboratories, Inc. ("Barr"), by its counsel, will take the deposition upon oral examination of GlaxoSmithKline.

The deposition will take place at Esquire Philadelphia, 1600 JFK Boulevard, Suite 1210, Philadelphia, Pennsylvania 19103 commencing on July 6, 2007 at 1:00 p.m, or at such other time and place as may be agreed to by counsel. The deposition will be taken before a Notary Public or other person authorized to administer oaths by the laws of the United States, and will be recorded by stenographic and/or videographic means. The deposition will continue from day-to-day until completed.

All counsel are invited to attend and cross-examine.

Pursuant to Rule 30(b)(6), GlaxoSmithKline shall designate in writing to Barr the names of the persons with knowledge or information reasonably available to GlaxoSmithKline on the matters set forth in Exhibit A attached hereto (subject to the Definition and Instructions

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indicated) who will testify on its behalf with respect to those matters, specifying the matters as to which that person will testify.

Dated: June 15, 2007

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EXHIBIT A

DEFINITIONS AND INSTRUCTIONS

The definitions and instructions set forth in the subpoena served on GlaxoSmithKline ("GSK") on August 3, 2006 in the above-captioned case shall apply and are incorporated herein by reference.

TOPICS FOR DEPOSITION

- Documents produced in response to the subpoena served on GSK on August 3, 1. 2006 in the above-captioned case.
- 2. The formulation for Flonase® nasal spray (NDA No. 020121) as of the approval date of NDA No. 020121, October 19, 1994, and any and all modifications to the Flonase® formulation from October 19, 1994 to the present and the date of any such modification.
- The formulation for Beconase AQ® nasal spray (NDA No. 019389) as of the 3. approval date of NDA No. 019389, July 27, 1987, and any and all modifications to the Beconase AQ® formulation from June 27, 1987 to the present and the date of any such modification.
- The size of the solid particles of fluticasone propionate medicament contained in 4. Flonase® nasal spray from October 19, 1994 to the present.
- The size of the solid particles of beclomethasone dipropionate monohydrate 5. medicament contained in Beconase AQ® nasal spray from July 27, 1987 to the present.

CERTIFICATE OF SERVICE

I, Karen E. Keller, Esquire, hereby certify that on June 15, 2007, I caused to be electronically filed a true and correct copy of the foregoing document with the Clerk of the Court using CM/ECF, which will send notification that such filing is available for viewing and downloading to the following counsel of record:

Steven J. Balick, Esquire John G. Day, Esquire Tiffany Geyer Lydon, Esquire ASHBY & GEDDES 500 Delaware Avenue, 8th Floor Wilmington, DE 19801

I further certify that on June 15, 2007, I caused a copy of the foregoing document to be served by hand delivery on the above-listed counsel of record and on the following non-registered participants in the manner indicated:

BY E-MAIL

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